

## DESCRIPTION

INHIBITOR OF CONSTITUTIVE ACTIVE FORMING OF CALCINEURIN

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## TECHNICAL FIELD

The present invention relates to an inhibitor of constitutive active forming of calcineurin. The present invention more specifically relates to a drug which inhibits cleavage of calcineurin subunit A (CaNA) by calpain.

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## BACKGROUND ART

Calcineurin is a phosphatase which is activated depending on calcium and calmodulin. It is a complex consisting of calcineurin subunit A (CaNA) having a catalytic center and calcineurin subunit B (CaNB) being a regulating factor. Calcineurin usually exists in an unactive form in the cell, because the association of the catalytic center with a substrate is obstructed by the autoinhibitory domain. The increased intracellular concentration of calcium leads to open the autoinhibitory domain by changing the structure of calcineurin (active form) which results in that the substrate can be associated with the catalytic center. Meanwhile, the decreased intracellular concentration of calcium leads calcineurin to return to the unactive form again. As just described, it is known in the prior art that calcineurin is reversibly activated depending on the concentration of calcium.

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In 1999, it was reported that calcineurin played an important role in neuronal cell death (refer to Asai Akio, et al., J. Biological Chemistry, Vol. 274. p. 34450, 1999). Furthermore, it has been also

reported that the calcineurin-specific inhibitor (immunosuppressants; FK506 and cyclosporin A) suppresses neuronal cell death (refer to Morioka Motohiro, et al., Progress in Neurobiology, Vol. 58, p. 1, 1999; and Springer Joe E., et al., The Journal of Neuroscience, Vol. 20, p. 7246, 5 2000). On the basis of many reports including such literatures, as a mechanism of neuronal cell death observed in brain ischemia and spinal injury, it has been suggested now that N-methyl-D-aspartic acid receptor (NMDA receptor), which is one of the glutamic acid receptors, is abnormally activated by the excitation of the neuronal cell followed by an 10 influx of large quantity of calcium into the cell through the receptor, and thereby calcineurin is activated to induce cell death. However, since the influx of calcium is transient, the activated state of calcineurin does not continue for a long term. Therefore, although it is imaged that there is any other mechanism by which calcineurin is activated for the long 15 term than the transient increase of intracellular concentration of calcium, such mechanism has never been reported.

As mentioned above, it is known that immunosuppressants FK506 and cyclosporine A have a suppressive effect on neuronal cell death, in particular ischemic and excitatory neuronal cell death. 20 However, it is known that the side effect of such agents is large (for example, toxicity, diabetic and the like), because the agents suppress all signal transduction involved in calcineurin. Therefore, a suppressant for neuronal cell death which can inhibit a long term activation of calcineurin and show less side effect compared with the conventional 25 one is strongly desired.

## DISCLOSURE OF INVENTION

The objects of the present invention are to solve the above-mentioned problems and to provide a suppressant for neuronal cell death having less side effect and effective for various diseases.

5 As a result of intensive investigations to solve the above-mentioned problems, the following findings were obtained.

a) When neuronal cell death was induced with administering glutamic acid, the fragmentation of calcineurin subunit A (CaNA) was observed.

10 b) Cleavage sites of CaNA were between amino acid residues of 392 (arginine) and 393 (lysine) and between amino acid residues of 421 and 425 in the amino acid sequence.

c) It was clarified that calcineurin had an activity independently of calcium and calmodulin, when it was cleaved at said 15 sites.

d) Said fragmentation was caused by calpain, which had thought to belong to a different signal transduction route from calcineurin, and inhibited by a calpain inhibitor.

e) When the peptide including amino acid residues 392-393 20 or amino acid residues 421-425 of CaNA was introduced in the neuronal cell, neuronal cell death by the glutamic acid administration was suppressed.

On the basis of the above-mentioned findings we have discovered a cell death suppressant targeting the mechanism of the long 25 term activation of calcineurin but not the calcineurin activation through the transient increase of the intracellular concentration of calcium, and have established the present invention. That is, this invention relates

to;

- (a) an inhibitor for cleavage of CaNA by calpain, comprising the peptide of SEQ No. 1, the peptide of SEQ No. 2 and/or analogues thereof;
- (b) a suppressant for neuronal cell death comprising the inhibitor for cleavage of CaNA of the above (a) as an active ingredient;
- (c) a suppressant for progress of a disease associated with dementia comprising the inhibitor for cleavage of CaNA of the above (a) as an active ingredient; and
- (d) an additive for a culture medium of cells and brain slice comprising the inhibitor for cleavage of CaNA of the above (a).

#### BRIEF DESCRIPTION OF DRAWINGS

Fig. 1 is photographs showing cleavage of CaNA on inducing neuronal cell death with the addition of glutamic acid, and the inhibitory effect of the CaNA inhibitor of the present invention on cleavage of CaNA.

Fig. 1 (a) shows the result of the sample which was incubated for 3 hours after the addition of glutamic acid, and Fig. 1 (b) shows the result of the sample which was incubated for 24 hours after the addition of glutamic acid. In Figs. 1 (a) and (b), lane 1 shows a control, lane 2 the sample in which glutamic acid was added alone, lane 3 the sample in which the cleavage inhibiting peptide was added alone, and lane 4 the sample in which glutamic acid and the cleavage inhibiting peptide were added.

Fig. 2 shows the number of neuronal cells in which neuronal cell death was induced by adding glutamic acid.

#### BEST MODE FOR CARRYING OUT THE INVENTION

The inhibitor for cleavage of calcineurin subunit A (CaNA:

SEQ No. 3) by calpain means a substance comprising the peptide of SEQ No. 1, the peptide of SEQ No. 2 and/or an analogue thereof, which inhibits cleavage of CaNA at between amino acid residues 392 and 393 or 421 and 425, and resulting in the inhibition of constitutive active 5 forming of calcineurin.

The above-mentioned analogue of the peptide of SEQ No. 1 includes a peptide comprising amino acid sequence of the peptide of SEQ No. 1 which is deleted, substituted a part thereof and/or another amino acid sequence is inserted or added at the terminal of the peptide, 10 and which inhibits cleavage of CaNA by calpain. For example, a peptide changed the amino acid residue 9 Arg to Lys and/or a peptide changed the amino acid residue 10 Lys to Arg in the peptide of SEQ No. 1 are included in the analogue of the peptide of SEQ No. 1 of the present invention.

15 The above-mentioned analogue of the peptide of SEQ No. 2 includes a peptide comprising amino acid sequence of the peptide of SEQ No. 2 which is deleted, substituted a part thereof and/or another amino acid sequence is inserted or added at the terminal of the peptide, which inhibits cleavage of CaNA by calpain. For example, a peptide 20 changed the amino acid residue 11 Lys to Arg in the peptide of SEQ No. 2 are included in the analogue of the peptide of SEQ No. 2 of the present invention.

An inhibitory activity of the analogue of the peptide of SEQ No. 1 or 2 (analogue peptide) for cleavage of CaNA can be evaluated by the 25 following manner. That is, the test solution (0.1  $\mu$ M or 10  $\mu$ M of analogue peptide, 5  $\mu$ M purified calpain, 20 mM Tris-HCl (pH 7.4) and 1 mM  $\text{CaCl}_2$ ) is prepared and then purified calcineurin is added thereto

and incubated at 30°C for 1 hour. The reaction solution is electrophoresed on a gel with 10 % SDS-PAGE, and is stained with Coomassie blue. The evaluation can be performed by determining the quantity of calpain dependent fragmentated calcineurins which are 5 found as molecular weights of 45 kDa and 48 kDa as the result of the stain.

The above-mentioned peptides can be prepared by using a known method such as solid phase or liquid phase synthesis with Boc or Fmoc strategy. The obtained peptide after that manner can be also 10 purified with a known method such as ether precipitation/filtration, high performance liquid chromatography (HPLC) or perfusion chromatography.

The inhibitor for cleavage of CaNA by calpain of the present invention can contain an additional compound as long as it contains the 15 peptide of SEQ No. 1, the peptide of SEQ No. 2 and/or the analogue thereof. The additional compound includes, for example, an intracellular transporting signal peptide consisting of 7 to 30 amino acid residues which contains 50 % or more arginine or lysine thereof, such as a polyarginine peptide (for example, polyarginine peptide consisting of 20 five arginine residues), protein transporting domain (PTD; SEQ No. 4) consisting of 11 amino acid residues which is contained in TAT protein of HIV virus; or liner polyethylenimine (PEI) being a cationic water-soluble polymer. Such compound can be associated with (or fused to) the peptide of SEQ No. 1, the peptide of SEQ No. 2 and/or the analogue 25 thereof by using a method such as (i) to be synthesized starting with the peptide of SEQ No. 1, the peptide of SEQ No. 2 and/or the analogue thereof using general peptide synthesis, or (ii) one peptide is bound to a

divalent crosslinking agent, a terminal of another peptide is bound to a cysteine residue, and then the both peptide is reacted. The thus obtained inhibitor for cleavage of CaNA can be purified with the above-mentioned purification method.

5        The suppressant for neuronal cell death of the present invention means an agent which comprises the above-mentioned inhibitor for cleavage of CaNA by calpain as an active ingredient, and inhibits long term induction of neuronal cell death in neuronal cells. The suppressant for neuronal cell death can be also used as an agent for  
10      preventing or treating a disease related to neuronal cell death. The suppressant for neuronal cell death of the present invention preferably contains the inhibitor for cleavage of CaNA comprising the peptide of SEQ No. 2 and the intracellular transporting signal peptide, and more preferably contains the inhibitor for cleavage of CaNA comprising the  
15      peptide of SEQ No. 1, the peptide of SEQ No. 2 and an intracellular transporting signal peptide. The above-mentioned agent for preventing or treating a disease related to neuronal cell death means a pharmaceutical preparation comprising an effective amount of the suppressant for neuronal cell death for preventing or treating the  
20      disease related to neuronal cell death. The disease related to neuronal cell death includes, for example, Alzheimer's disease, dementia disease, brain ischemic disease, intracerebral hemorrhage such as subarachnoid hemorrhage, spinal injury (trauma), Parkinson disease, epilepsys, and the like.

25        As an administration route of the suppressant for neuronal cell death of the present invention, there include, for example, oral administration, intravenous administration, intracerebral directly

administration and the like. The oral administration is more preferable in the light of a burden on a patient and the side effect.

A dosage form of the suppressant for neuronal cell death of the present invention can be appropriately set depending on the 5 administering method. Concretely, there include liquid formulations such as solution, emulsion and suspension, tablet, capsule, and the like. For example, in case of oral administration, tablet or capsule formulation is preferable, in case of intravenous administration or intracerebral directly administration, the liquid formulation is preferable. 10 Various additives usually used by a person skilled in the art according to the dosage form can be used for formulating the suppressant for neuronal cell death of the present invention. Such additives include an antioxidant, pH adjustment, preservative and the like.

The dosage of the above-mentioned suppressant for neuronal 15 cell death can be appropriately set depending on the administration method, age, weight and condition of the patient to be applied, and the like. For instance, the dosage is preferably at least 0.1 mg/kg/day as the inhibitor for cleavage of the CaNA by calpain of the present invention, and more preferably at least 1 mg/kg/day. When the dosage is less 20 than 0.1 mg/kg, it tends to reduce the inhibitory effect on cleavage of CaNA by half. Moreover, the dosage is preferably 100 mg/kg/day or less as the inhibitor for cleavage of the CaNA by calpain of the present invention, and more preferably 20 mg/kg/day or less. When the dosage exceeds 100 mg/kg, it tends to show the cell toxicity. The 25 suppressant for neuronal cell death of the present invention can be administered by either single dose or multiple doses.

An additive for a culture medium of cells and brain slice

comprising of the present invention means the additive for the culture medium containing at least the inhibitor for cleavage of CaNA. The additive for the culture medium of cell and brain slice comprising of the present invention preferably contains the inhibitor for cleavage of CaNA  
5 comprising the peptide of SEQ No. 2 and the intracellular transporting signal peptide, and more preferably contains the inhibitor for cleavage of CaNA comprising the peptide of SEQ No. 1, the peptide of SEQ No. 2 and an intracellular transporting signal peptide.

When the inhibitor for cleavage of CaNA by calpain of the  
10 present invention is applied to culture cells, the amount of addition is preferably 0.01 to 100 nmol/ml, and more preferably 0.1 to 10 nmol/ml in the culture medium having the cell concentration of  $1 \times 10^5$  cell/ml. When the amount of addition is less than 0.01 nmol/ml, it tends to reduce the inhibitory effect on cleavage of CaNA by half, and when it is  
15 more than 100 nmol/ml, it tends to show the cell toxicity.

The present invention is explained in more detail by means of the following Examples, but the present invention is not limited thereto.

#### EXAMPLE 1

20 In order to introduce FDGATAAARKEVIRNK (SEQ No. 1) and REESESVLTLKGLTPTG (SEQ No. 2) as the inhibitor for cleavage of CaNA of the present invention into the cultured cells, the following oligopeptides in which the intracellular introducing signal peptide (ten arginines) was added to their N terminal were prepared (available from  
25 PEPTIDE INSTITUTE INC.).

RRRRRRRRRRFDGATAAARKEVIRNK (SEQ No. 5)

RRRRRRRRRRREESESVLTLKGLTPTG (SEQ No. 6)

(Preparation of neuronal cells)

After the brain hippocampus from 18 days embryo of Wister rat had been removed, it was treated with PBS containing 0.05 % trypsin for 15 minutes at 37°C. After the neuronal cells were 5 suspended with a glass pipet,  $1 \times 10^6$  cells were cultured in 3.5 cm in the diameter culture dish which was previously coated with poly-D-lysine. As the medium, 3 ml Neuro Basal medium (available form Invitrogen, Inc.) complemented with B27 supplement (0.03 ml; available from Invitrogen, Inc.), penicillin (100 units/ml in the final 10 concentration; available form Invitrogen, Inc.) and streptomycin (100 µg/ml in the final concentration; available form Invitrogen, Inc.) was used and the cultivation was carried out in carbon dioxide incubator (5 % CO<sub>2</sub>, 37°C).

15 (Addition of peptide)

The peptides of above-mentioned SEQ Nos. 5 and 6 were added in the culture solution with the final concentration of 1 µM 10 days after beginning of culture, and it was incubated in carbon dioxide incubator (5 % CO<sub>2</sub>, 37°C). Three hours after the addition, glutamic 20 acid was added in the final concentration of 500 µM, and it was incubated for 15 minutes. The culture solution was then exchanged, and it was further cultured. The cells were collected at 3 hours and 24 hours after glutamic acid was added, were disrupted by ultrasonic wave in 1 % SDS solution, and the SDS-PAGE buffer was added thereto. 25 After this sample was subjected to the SDS-PAGE gel electrophoresis, Western blotting was carried out using the antibody recognizing CaNA (rabbit serum, available form Santa Cruz Biotechnology, Inc). In this

Example, a cell sample in which neither glutamic acid nor the cleavage inhibiting peptide was added was used as a control. Moreover, the cell sample in which only glutamic acid was added and the cell sample in which only the cleavage inhibiting peptide was added were concurrently prepared.

The result was shown in Fig. 1. In the glutamic acid addition group, the fragmentation of calcineurin was observed. On the other hand, in the neuronal cells treated with the cleavage inhibiting peptide, cleavage of calcineurin which was able usually to be caused by the addition of glutamic acid was inhibited.

## EXAMPLE 2

The neuronal cells were cultured in the same manner as in Example 1, and then the above-mentioned peptides of SEQ Nos. 5 and 6 were added in the final concentration of 1  $\mu$ M in the same manner as in Example 1. The calcineurin inhibitor, FK506 (available from Fujisawa pharmaceutical Co., Ltd.; the final concentration of 1  $\mu$ M) or calpain inhibitor, ALLM (available from Merck & Co., Inc.; the final concentration of 25  $\mu$ M) was added as a control. After 2 hours incubation with each reagent, glutamic acid was added in the final concentration of 500  $\mu$ M, followed by 15 minutes incubation. The culture solution was then exchanged, and it was further cultured. Each neuronal cell was fixed with 4 % paraformaldehyde 3, 6, 12 or 24 hours after the addition. Thereafter, TUNEL staining (available from Roche Diagnostics, Inc) was carried out to identify the neuronal cells which caused cell death. The TUNEL positive cell was counted.

The result is shown in Fig. 2. In the glutamic acid addition

group, the number of the neural cells which caused cell death increased with the time course after the administration. It was demonstrated that the cleavage inhibiting peptide had the suppressive effect on neuronal cell death induced by glutamic acid. Moreover, the effect was the same 5 degree as that of FK506 or ALLM.

#### REFERENCE EXAMPLE 1

Calcineurin 1  $\mu$ M refined from cow brain was reacted with 1  $\mu$ M of calmodulin (available from Merck & Co., Inc.) and/or 1  $\mu$ M of m-10 calpain (available from Merck & Co., Inc.) in a reacting solution (containing 20 mM Tris-HCl, pH 7.4, 1 mM  $\text{CaCl}_2$ , and 1 mM  $\text{MgCl}_2$ ) at 30°C for one hour, respectively. After the reaction 12 % SDS-PAGE gel electrophoresis was carried out and then said gel was stained with Coomassie blue.

15 As a result, in case that the calpain was not added, the purified CaNA was seen at the size of 60 kDa on the electrophoresis. This is corresponding to the size of CaNA that has been reported up to now. On the other hand, in case that calmodulin and calpain were added, the band was not observed at 60 kDa but was observed at 48 and 20 45 kDa. In case that calcineurin was reacted with only calpain, the cleaved CaNA in the size of 45 kDa was observed.

As the above-mentioned result it is clear that calcineurin is cleaved by calpain.

In addition, as determining the cleavage site of CaNA by 25 calpain, the cleaved protein of 45 kDa was composed of the amino acids up to 392 in the amino acid sequence and the cleaved protein of 48 kDa was composed of amino acids up to 421, up to 422, up to 423 and up to

424 in the amino acid sequence.

#### INDUSTRIAL APPLICABILITY

This invention provided the inhibitor for cleavage of CaNA.

5 The inhibitor for cleavage of CaNA of the present invention can suppresses neuronal cell death caused by the activation of calcineurin, because it can inhibit the constitutive active forming of calcineurin. Moreover, a suppressant for neuronal cell death of the present invention containing the inhibitor for cleavage of CaNA is excellent useful because  
10 it can be used as a drug for prevention or treatment of the disease associated with neuronal cell death including the dementia disease and the like. In addition, if the additive of the culture medium for cells of the present invention containing the inhibitor for cleavage of CaNA is used, the cultured cell can be grown well. Moreover, the inhibitor for  
15 cleavage of CaNA of the present invention can be also used as a reagent for the investigation about neuronal cell death.

#### SEQUENCE LISTING FREE TEXT

SEQ No. 5: Peptide sequence consisting of peptide sequence derived  
20 from human and artificial peptide sequence.

SEQ No. 6: Peptide sequence consisting of peptide sequence derived from human and artificial peptide sequence.